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BADI 92.09.21 A(8-N2) C(7-H1, 10-A12C, 14-A1, 14-A184, 14-A4) D(0-H1, 1 7-B, 9-A1C) E(7-A1, 7-B1, 7-B2, 7-C, 7-D4B, 7-D8, 7-F1, 10-A18A) G(3-A) H(7-G)	*DE 4231518-A1 trl-substd. by 1-4C alkyl, 1-4C haloalkyl, 1-4C alkylthio or 1-4C alkoxy, 1-4C alkylthio or 1-4C haloalkylthio); A6, 307/68, A = a cyclic gp. of formula (A1) to (A7):		8. of (R ³)n Ty Me	(A1) (A2) (A3) (A4)	$(R^3)_n$ R^4 R^4 R^6	3-12C	ycloalkyl, (A5) (A6) (A7)	no- to
94-102435713 A60 C03 D22 E14 G06 H H07 (C02 D17 D18 D21 E13)	BASF AG 92.09.21 92DE-4231518 (94.03.24) C07D 213/82, A01N 37/28, 43/16, 43/32, C07D 277/56, 309/28, 333/38, 327/06, 307/68, 231/14, A01N 43/40, C07C 259/10	N-Hydroxy-N-phenyl-carboxyllc acid amide derivs for use as microbicides, esp. fungicides, and are partic. effective against Botrytis	N-hydroxy-N-phenyl carboxylic acid amide derive. of formula (I) are new:	€	R	R = 2-12C alkyl, 2-12 C alkoxy, 3-12C alkenyl, 3-12C alkenyloxy, 3-6C alkynyl, 3-6C alkynyloxy (all opt	partially or completely halogenated), 3-7C cycloalkyl, 4-7C cycloalkyloxy or 4-7C cycloalkyloxy or 4-7C cycloalkyloxy or 4-7C cycloalkyloxy or 4-7C	or tri substd. by 1-4C sikyl), Ph (opt. mono- to

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V

were added at 0°C to a soln. of 15.1g 2-n-propyl phenyl hydroxylamine in 75 ml ether-petroleum ether mixt. (2:1); followed by dropwise addn. of 13.6g 2-chloronicotinic acid chloride with intensive mixing. After stirring the mixt. overnight at room temp., the residue was filtered off and worked up to give 14.6g amide prod. which was	recrystd. from OH to give 12.5g 2-chloronicotinic acid-N-hydroxy-2-n-propylanilide of m.pt. 134-135°C. (26pp14010BDwgNo0/0). Addnl. Doig: EICKEN K, AMMERMANN E, LORENZ G	, DB4231518-A
$X = CH_2$, S, SO or SO ₂ ; Y = 0 or S; R^1 , R^2 , R^4 , R^5 and R^7 = halogen, 1-4C alkyl or 1-4C haloalkyl; R^3 , R^6 = H, halogen or 1-4C alkyl; R = 1-2.	OSE (1) are fungledes for control of harmful fungi, esp. of Botrytis. They can be used to protect materials, e.g. glues, starch solns, leather, lubricants, etc. or plants from attack by bacteria or fungi. (1) are effective against e.g. Staphylococcus aureus, Escherichia coli, Saccheromyces cereviscise, Candida alticans, etc. PREPARATION	EXAMPLE Ho-NH

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N-hydroxy-N-phenylcarboxamides, processes for their production and agents containing them for the controlling of harmful fungi.

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Inventor:

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Applicant:

BASF AG (DE)

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C09D5/14; C09G1/08; C09H5/00; C09J11/06; C09K3/10; C09K7/02; C10L1/00; C10M133/16; C10M133/38; C10M135/32; C14C1/00; C14C9/00; D06M13/402; D21F1/66; C07D213/82; A01N37/28; A01N43/16; A01N43/32; A01N43/40; C07C259/10; C07D231/14; C07D277/56; C07D307/68; C07D309/28; C07D327/06; C07D333/38; C08K5/20; C08K5/34;

C08K5/47

- european:

A01N43/40; A01N43/78; C07D213/82G; C07D231/14;

C07D277/56; C07D307/68; C07D309/28; C07D327/06;

C07D333/38

Application number: DE19924231518 19920921
Priority number(s): DE19924231518 19920921

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Abstract not available for DE4231518
Abstract of corresponding document: **EP0591699**

1. N-hydroxy-N-phenylcarboxamides of the formula I in which the substituents have the following meaning: R is optionally substituted alkyl, alkoxy, alkenyl, alkenyloxy, alkinyl, alkinyloxy, cycloalkyl, cycloalkenyl, cycloalkyloxy, cycloalkenyloxy or phenyl; A is one of the radicals A1 to A7 where X is -CH2-, -S-, -SO- or -SO2-; Y is -O- or -S-; R<1>, R<2>, R<4>, R<5> and R<7> are halogen, alkyl or haloalkyl; R<3> and R<6> are hydrogen, halogen or alkyl; n is 1 or 2; their preparation, agents containing them, and their use for controlling harmful fungi.

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